

MORBIDITY AND MORTALITY WEEKLY REPORT

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Importation of Wild Poliovirus into Qinghai Province — China, 1999

Indigenous wild poliovirus was last isolated in China in 1994. On October 13, 1999, a case of acute flaccid paralysis (AFP) in a 16-month-old boy was reported to public health authorities in Xunhua Autonomous County, Haidong Prefecture, Qinghai Province, China. Following onset of paralysis on October 12, the boy was no longer able to stand or walk. Two stool samples, taken within 14 days of onset of paralysis, were analyzed in the Qinghai provincial laboratory and yielded poliovirus. The isolates were later differentiated as wild poliovirus type 1 at the National Poliovirus Laboratory in Beijing. Stool specimens from one of five children with whom the boy had contact yielded wild poliovirus type 1. This report describes this case of poliomyelitis and the public health response to the case in China.

The case occurred among the Sala, a group of approximately 80,000 persons who live mainly in Xunhua Autonomous County, Qinghai, or in neighboring Gansu province. Many Sala are traders, and Sala men travel widely within Qinghai and to nearby provinces, including Gansu, Sichuan, and Xinjiang, and to Tibet as far south as the border with Nepal. The Sala have trade contacts in India, Pakistan, and Central Asia. Neither the casepatient nor immediate family members are reported to have traveled outside Xunhua County during the 2 months before paralysis onset.

Despite intensive investigations, including retrospective record reviews in health-care facilities and active case searches in villages in selected areas, no additional polio cases or other evidence of continued poliovirus circulation was found. Since 1996, the quality of AFP surveillance in Qinghai has been excellent, with nonpolio AFP rates of >1.5 per 100,000 population and proportion of cases with two adequate stool specimens between 70%–90% annually. The provincial laboratory in Qinghai has shown proficiency in 1999 and received full accreditation within the World Health Organization polio laboratory network.

The Qinghai poliovirus strain is closely related (98%) to poliovirus isolates from central and northern India during 1998–1999, but unrelated to polioviruses that circulated in China until 1994. Despite the absence of a history of travel by the case-patient or his immediate family, evidence suggests that the virus was imported from a neighboring country, probably India, where polio is endemic. The extent of virus circulation following importation has not yet been determined (the paralytic case-to-infection ratio is typically 1:200 in a fully susceptible population). No evidence exists of continued circulation of poliovirus.

Importation of Wild Poliovirus into Qinghai Province - Continued

Before confirmation of the index case (but after onset of paralysis), provincewide supplementary vaccination with oral poliovirus vaccine, planned earlier in 1999 and targeting children aged 0–3 years, was carried out in late November in both Qinghai and Tibet. In response to confirmation of the index case, an initial local case-response vaccination round was conducted in Xunhua County in November. This was followed by round 1 of a larger, intense house-to-house mopping-up vaccination activity targeting children aged 0–9 years that was implemented in six of eight prefectures of Qinghai, beginning in early December. Round 2 in January 2000 also included house-to-house mopping-up vaccination targeting 7.1 million children in an even larger area, including Qinghai, Ningxia, most of Gansu, and parts of Tibet. These extensive mopping-up vaccination activities were in addition to the second round of subnational immunization days conducted January 5–6, 2000, in all provinces in high-risk areas to vaccinate children aged 0–3 years. All vaccination activities reported good coverage of the target population. Two additional large multiple-province vaccination rounds, targeting approximately 26 million children, are planned for March and April.

Since the case was identified, surveillance activities have been intensified through active case searches in health-care facilities and communities during mopping-up vaccination and retrospective review of hospital records. Special assessments of the quality of virologic surveillance were conducted, including specimen collection and handling procedures, and the quality of specimen processing at the provincial laboratory.

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Editorial Note: Preliminary data from this investigation suggest that the polio case in Qinghai was caused by importation of wild poliovirus with limited circulation. No other cases have been detected despite high-quality AFP surveillance and extensive searches of hospital records, health-care facilities, and communities. Further intensive surveillance and vaccination activities, including active house-to-house searches for recent AFP cases, are being conducted.

The detection of this case in a sparsely populated rural area of China indicates that high-quality AFP surveillance continues to be maintained in China. The detection also highlights the need for all polio-free countries to remain vigilant to allow early detection of wild poliovirus imported from countries where polio is endemic and to institute rapid control measures.

Role of Victims' Services in Improving Intimate Partner Violence Screening by Trained Maternal and Child Health-Care Providers — Boston, Massachusetts, 1994–1995

From 1992 to 1996, approximately 1 million incidents of nonfatal intimate partner violence (IPV) occurred each year in the United States; 85% of victims were women (1). In 1989, pediatric research found a concurrence of victimization of mothers and their children and supported a recommendation that maternal and child health-care providers (HCPs) pursue training and advocate for increased access to services to promote the safety and well-being of mothers and their children (2). From 1992 to 1997, the Pediatric

Intimate Partner Violence Screening - Continued

Family Violence Awareness Project (PFVAP), a training project for maternal and child HCPs, promoted prevention of and intervention for IPV in Massachusetts (3). In 1994, PFVAP conducted a pilot evaluation in two urban community health centers to determine whether HCPs trained to conduct IPV assessment would increase their screening rates of women at risk for IPV if an on-site referral service for victims was available. This report summarizes the results of the pilot project, which indicate that IPV screening rates did not increase after implementing on-site victim services.

Screening rates were assessed for 14 HCPs at two community health centers (centers A and B) in a low-income, racially mixed, urban community in the Boston area. Because the two centers were dissimilar in patient demographics and other characteristics, one could not be compared with the other. Therefore, a phased intervention design was used; IPV screening was measured during two 10-week periods (phases 1 and 2). Phase 1 followed a 2-hour group training session to teach HCPs to implement a brief screening protocol* of female patients and mothers of pediatric patients aged 0–12 years during routine visits using a recommended screening schedule.† Phase 2 followed implementation of on-site victim services that offered weekly support groups separately for battered women and children using the identical protocol as in Phase 1. Between the end of phase 1 and the beginning of phase 2, there was a 3-month period.

To document screening in each phase, HCPs recorded during each visit with each female adult patient and each mother of a pediatric patient whether 1) the patient received IPV screening and who performed the screening; 2) any family members were present during the patient visit; and 3) a staff interpreter was present during the visit. Date of birth, race/ethnicity, marital status, date and type of visit, and diagnosis were gathered from the patients' files. A physician subsequently coded diagnoses into the following categories: routine health-care maintenance, prenatal care, acute/sick, chronic problem, injury, psychosocial, human immunodeficiency virus/sexually transmitted diseases (HIV/STD), and pain.

For both phases, an observed screening rate was calculated for each HCP and defined as the proportion of the HCPs' patients seen and screened by the HCP during that period. Although the PFVAP protocol recommended screening some patients (pregnant women and mothers of children aged <2 years) more than once a year, patients who were screened at least once during phase 1 were considered "previously screened" and were not included in calculating phase 2 screening rates.

The combined data from both health centers and both phases (after exclusions) (Table 1) comprised 14 HCPs, 642 patients, and 1352 patient visits. Each patient's final screening status (ever or never screened) was based on combined data from each phase and was evaluated relative to patient demographics and visit characteristics by two separate logistic regression models.

*Suggested questions were 1) "I ask all my patients, do you feel safe in your home?"; 2) "Is anyone hurting you, harassing you, or making you feel afraid?"; and 3) "At any time, has your partner ever pushed, hit, or kicked you?"

The recommended schedule consisted of screening 1) adult and adolescent females during routine gynecologic, internal or family medicine, or pediatric visits annually; 2) mothers of pediatric patients aged 2–12 years annually; 3) mothers of pediatric patients aged 0–2 years twice annually; and 4) patients during prenatal-care visits once per trimester.

Intimate Partner Violence Screening - Continued

TABLE 1. Inclusion and exclusion criteria for health-care providers (HCPs) and patient visits for intimate partner violence (IPV) screening — Boston, Massachusetts, 1994–1995

Level	Inclusion criteria	Exclusion criteria
HCPs	Met with ≥26 patients during study period	Met with ≤25 patients during study period
Patient visits	Scheduled at least 1 day in advance	Visits by females aged 13-17 years*
	"Screening target" [†] present	Adult other than screening target in room with HCP ⁵
		For phase 2: patients screened during phase 1

*Excluded because two possible screening targets (the mother or the adolescent female) could have been in the room with the HCP. HCPs' documentation of screening was unclear about whether mothers or adolescent females were interviewed for IPV risk.

¹ A woman aged ≥18 years or the female caretaker of a pediatric patient aged 0-12 years.

For the safety of patients and HCPs, HCPs were instructed not to screen for IPV risk if adults other than the screening target and a staff interpreter were in the room.

Source: Pediatric Family Violence Awareness Project Evaluation

Eleven (79%) of 14 HCPs did not demonstrate increased screening during phase 2, following on-site services implementation. Unadjusted combined screening rates for both health centers decreased significantly from phase 1 (33% patients screened) to phase 2 (23%) (p<0.03). For each phase, health center A had approximately twice the documented screening rate of health center B. On average, screening rates declined 7.4% (standard deviation [SD]=15.7 percentage points) at health center A and 14.1% (SD=17.5 percentage points) at health center B.

At both health centers, unadjusted individual HCP screening rates varied during both phases from 1.8% to 92.8% during phase 1 and from 0 to 94.9% during phase 2. The degree of change in HCP screening rates also varied widely. Individual HCP screening rates of decline ranged from 1.8 to 46.6 percentage points. For the three HCPs who demonstrated increases between phase 1 and phase 2, the increase ranged from 0.6 to 24.7 percentage points.

Analyses of visit, HCP, and patient characteristics controlled for health center and used combined rates from both phases to improve the stability of estimates. Several aspects of patient visits predicted the likelihood of screening. Patients were screened more often during routine visits (p<0.01). However, screening was 23 times more likely during adult medical visits (p<0.01) and 10 times more likely during gynecologic visits (p<0.01) than during pediatric visits. Diagnostic categories also were related significantly to screening status. Patients seeking treatment for pain were four times more likely to be screened (p<0.03). A combined variable of injury, HIV/STD, and psychosocial problems also was a significant predictor of screening (p<0.04). Of the patient characteristics examined, only unknown marital status was a significant predictor of screening status, with women of unknown marital status less likely (p<0.01) to be screened than married patients.

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Intimate Partner Violence Screening - Continued

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Editorial Note: The results of this study suggest that the availability of on-site services for IPV victims alone may not be sufficient to overcome trained HCPs' perceptions of IPV as a problem for which they are ill-prepared to intervene (4). Systems approaches, such as continuous quality improvement in community health centers, may be more likely to sustain improved IPV screening rates through institutional policies linked to accountability (5). The impact of case mix on provider- and institutional-level IPV screening rates also requires more study. However, clinicians' adherence to the recommended practices to screen routinely all women at risk for IPV should be encouraged (6,7).

The findings in this report are subject to at least three limitations. First, because a convenience sample of community health centers was used, the results cannot be generalized to other community health centers or HCPs in the rest of Massachusetts or elsewhere. Second, the quasi-experimental design, which lacked a concurrent control, does not account for secular changes in screening behavior that may have occurred over the course of the study. Finally, phase 2 was delayed to involve the community health centers' administrative and clinical staff in the process of selecting IPV advocates and to address other administrative details of service development. Because data were not collected on the screening rates of HCPs immediately before phase 2, the effects of the on-site victims' services on individual HCPs cannot be determined fully.

Maternal and child HCPs see many battered women and their children in various settings, but rarely ask about family violence and IPV (6-9). Practitioners need additional training and support to assess and manage complex cases of family violence longitudinally (10). Further research to explore effective IPV interventions in health-care settings is needed.

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Information Needs and Uses of the Public Health Workforce — Washington, 1997–1998

Substantial efforts have been made to ensure that state and local public health agencies have the information technology and training needed for public health communications, information access, and data exchange (1,2). Numerous public health-related data and information resources are available on the World-Wide Web (e.g., MEDLINE, MMWR, CDC Prevention Guidelines Database, and Emerging Infectious Diseases); however, little systematic work has been done to understand the information needs of the public health workforce. To identify these needs and patterns of use and to set priorities for developing new online public health information resources, the University of Washington School of Public Health and Community Medicine (UW SPHCM) and the Washington State Department of Health (WSDOH) held structured and facilitated discussions with segments of the local public health workforce in Washington during 1997–1998. This report summarizes the results of those discussions, which indicate that different segments of the public health workforce have different information needs.

Five subgroups of the local public health workforce were selected for inclusion in the investigation on the basis of input from state and local public health leaders: 1) local health officers and public health agency directors, 2) environmental health directors, 3) directors of public health nursing, 4) health assessment coordinators and epidemiologists, and 5) a group comprising public health officials from small local health departments in which staff typically have responsibilities in multiple areas (e.g., nursing and disease investigation). Open-ended questions about information acquisition and use were developed in consultation with UW SPHCM faculty, WSDoH leaders, and staff from the Eastern and Western Washington Area Health Education Centers (AHECs). AHEC directors served as facilitators at each discussion.

Eight sessions were held from June 1997 through April 1998. A total of 70 persons participated; the smallest group had four and the largest had 14 participants. Persons in each group were from a cross section of local health jurisdictions representing metropolitan and rural areas, large and small agencies, and eastern and western Washington. The participants included 22 environmental health directors (in two sessions in different parts of the state), 10 public health nursing directors, 13 health assessment coordinators and epidemiologists (in two sessions in different parts of the state), four health officers/agency directors, and 21 staff members (mixed segments) from two small county health departments.

Seven information needs were identified by all four workforce segments (Table 1):

1) better tools and resources for contacting experts; 2) updates on pertinent legislative issues and events; 3) structured information ("metadata") characterizing the contents of data sets; 4) outcome measures and "best practice" resources; 5) better scheduling software and event calendars; 6) standard templates for frequently used applications; and 7) synthesized, knowledge-based information from external databases. Five needs were identified by more than one group and another 15 needs were identified by a single group (Table 1).

Interest in the use of information resources and technology also varied across groups (e.g., nursing directors expressed more interest in using videoconferencing technology than did other groups [Table 1]). Some groups expressed readiness to incorporate online resources (e.g., contact lists, statistical databases, and Web-accessible knowledge resources) into their work.

Information Needs and Uses of the Public Health Workforce - Continued

TABLE 1. Data and information resource needs of four local public health workforce segments — Washington, 1997–1998

	Assessment coordinators and pidemiologists	Nursing directors	Environmental health directors	Health officers and agency directors
Access to academic/state experts	Х	Х	×	Х
Administrative/budget data				X
Notification of continuing education opportunities			x	
Criminal justice data	X			
Disease incidence data (county/state/national)	×	X		
Disease/condition information*	×			
Geographically coded health-related data			X	
Health education information for the public			×	
Health education program information		X		
Health insurance billing data	X			
Vaccination guidelines		X		
Industrial effluent data			×	
Laboratory data (online)			×	
Laws and regulations (county/state)			×	X
Legislative issues updates	X	X	X	X
Local/small area data	X			
Metadata on data sets ¹	×	X	X	X
Outcome measurement resources	X	X	X	X
Group-specific electronic discussion list	s X		X	
Remote access to office systems and meetings		X		
Scheduling software/resources	×	X	X	X
Socioeconomic data	×			
Standard templates ¹	×	X	X	X
State agency data/resources/publication	s X	X		
Synthesized, knowledge-based informati	ion ¹ X	X	X	X
Treatment data**	×	X		
U.S. census data	X			

^{*}Includes fact sheets, nursing protocols, treatment for contacts, epidemiologic summaries, and prevention guidelines.

¹Include information on scope, coverage, location, how to access, and strengths and weaknesses of the data.

⁶E.g., reporting forms, surveys, assessment instruments, and management tools.

¹Include custom synthesized information and access to online bibliographic and factual databases (e.g., MEDLINE and CDC Prevention Guidelines Database).

^{**}Include hospital-based and clinic-based ambulatory, emergency, and inpatient care.

Information Needs and Uses of the Public Health Workforce - Continued

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Editorial Note: Public health practice spans numerous health, environmental, and social science disciplines; therefore, public health practitioners need access to diverse and complex information and data from multiple sources. Electronic access to peer-reviewed biomedical literature is available through MEDLINE (3); however, this resource meets only a portion of the public health practitioner's information needs (4). The variety in the types of information needed is matched by the diversity of the public health workforce itself that includes agency directors, environmental health scientists, epidemiologists and health assessment specialists, health educators, health officers, laboratorians, nurses, nutritionists, sanitarians, social workers, and outreach workers. Ideally, the development of online public health information resources should reflect this complexity and diversity.

Approximately one fourth of the information needs identified in this study was shared by all segments of the Washington public health workforce, but nearly half of the information needs was not shared by more than one segment. Also, readiness to incorporate the use of online information resources into public health practice varied across segments. In addition to diverse information needs, these findings may reflect differences in training, experience, and professional culture.

This study is subject to at least two limitations. First, these data are based on interviews with public health professionals in Washington only and may not represent the information needs in other states. Second, some public health workforce groups were not interviewed (e.g., health educators, nutritionists, social workers, and other outreach workers); therefore, the study probably underestimates the range and diversity of information needs among public health workers.

CDC's Information Network for Public Health Officials (1), the Health Alert Network (2), and the National Library of Medicine's Partnership in Information Access for Public Health Officials (5) are designed to strengthen the information infrastructure of state and local public health agencies. The success of these initiatives will depend not only on technology but also on the information content being delivered and used and on a workforce trained to use effectively these new tools and resources. Further research is needed to determine optimal development, structure, delivery, and marketing of public health information to specific public health workforce segments.

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Notice to Readers

Satellite Broadcast on Genital Dermatology

The National Network of STD/HIV Prevention Training Centers will present "STD Grand Rounds: Genital Dermatology," a national satellite broadcast on Thursday, March 9, 2000, from 1 to 3 p.m. eastern standard time. This program is for clinicians at sites across the United States and will be available in English or Spanish. The program is produced by the New York State Centers for STD/HIV Prevention Training in collaboration with the STD/HIV Prevention Training Center of New England. The broadcast is jointly sponsored for continuing medical education credit by the University of Cincinnati and for continuing education unit credit by the Massachusetts Department of Public Health.

Information on attending at a prearranged site or an alternate site is available from the STD/HIV Prevention Training Center in each public health region: Region I (Connecticut, Maine, Massachusetts, New Hampshire, Rhode Island, and Vermont), telephone (617) 983-6945; Region II (New Jersey, New York, Puerto Rico, and U.S. Virgin Islands), telephone (518) 474-1692; Region III (Delaware, District of Columbia, Maryland, Pennsylvania, Virginia, and West Virginia), telephone (410) 396-3876; Region IV (Alabama, Florida, Georgia, Kentucky, Mississippi, North Carolina, South Carolina, and Tennessee), telephone (205) 930-1154; Region V (Illinois, Indiana, Michigan, Minnesota, Ohio, and Wisconsin), telephone (513) 558-3197; Region VI (Arkansas, Louisiana, New Mexico, Oklahoma, and Texas), telephone (214) 819-1947; Region VII (Ilowa, Kansas, Missouri, and Nebraska), telephone (314) 747-0294; Region VIII (Colorado, Montana, North Dakota, South Dakota, Utah, and Wyoming), telephone (303) 436-7226; Region IX (Arizona, California, Hawaii, Nevada, and the Pacific Islands), telephone (510) 883-6600; and Region X (Alaska, Idaho, Oregon, and Washington), telephone (206) 685-9850. Registration also is available through the World-Wide Web at http://www.stdptc.uc.edu.**

Sites must be registered for participants to receive the handouts and continuing education credit. Additional information is available by telephone, (888) 232-3299 (or for persons with hearing impairment, [877] 232-1010); enter document number 130035 when prompted.

Notice to Readers

Availability of Draft of Updated Guidelines for Evaluating Surveillance Systems

A surveillance system enables ongoing collection, analysis, and dissemination of data to prevent and control disease or injury. Because all surveillance systems should be assessed periodically for their purpose and usefulness, in 1988 CDC published *Guidelines for Evaluating Surveillance Systems* (1). Recent developments in the electronic exchange of health data, the establishment of data-collection standards, and interest in

^{*}References to sites of non-CDC organizations on the World-Wide Web are provided as a service to MMWR readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

Notices to Readers - Continued

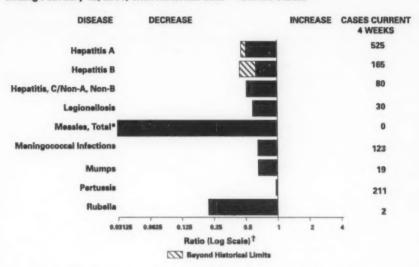
the integration of health information and surveillance systems have resulted in the need to update CDC's quidelines (2).

After researching and discussing various issues related to public health surveillance systems, the CDC Guidelines Working Group has composed a draft of *Updated Guidelines for Evaluating Surveillance Systems*. A copy of this draft is available on the World-Wide Web at http://www2.cdc.gov/revguide/index.htm (user name=community; password=guidelines) or by mailing a request for a copy to CDC Guidelines Working Group, Epidemiology Program Office, Mailstop K74, 4770 Buford Highway, Atlanta, GA 30341-3717. Comments about the draft of the updated guidelines should be submitted at the above Internet site or by mail by March 31, 2000.

References

- 1. CDC. Guidelines for evaluating surveillance systems. MMWR 1988;37(no. S-5).
- 2. CDC. Revising CDC's guidelines for evaluating surveillance systems. MMWR 1998;47:1083.

FIGURE I. Selected notifiable disease reports, comparison of provisional 4-week totals ending February 12, 2000, with historical data — United States



*No measles cases were reported for the current 4-week period, yielding a ratio for week 6 of zero (0).

Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary — provisional cases of selected notifiable diseases, United States, cumulative, week ending February 12, 2000 (6th Week)

		Cum. 2000		Cum. 2000
Anthrax			HIV infection, pediatric**	9
Brucellosis*		3	Plague	1
Cholera			Poliomyelitis, paralytic	
Congenital rubo	ella syndrome	1	Psittacosis*	
Cyclosporiasis*		2	Rabies, human	
Diphtheria			Rocky Mountain spotted fever (RMSF)	19
Encephalitis:	California® serogroup viral		Streptococcal disease, invasive Group A	301
	eastern equine*		Streptococcal toxic-shock syndrome®	16
	St. Louis®		Syphilis, congenital ¹	
	western equine®		Tetanus	
Ehrlichiosis	human granulocytic (HGE)*	4	Toxic-shock syndrome	13
	human monocytic (HME)*	1	Trichinosis	1
Hansen Disease	•	3	Typhoid fever	26
Hantavirus puli	monary syndrome*1		Yellowfever	
Hemolytic uren	nic syndrome, post-diarrheal®	6		

-: no reported cases

*Not notifiable in all states.

"Not notifiable in all states.

'Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID).

'Updated monthly from reports to the Division of HIV/AIDS Prevention—Surveillance and Epidemiology, National Center for HIV,

STD, and TB Prevention (NCHSTP), last update January 30, 2000.

'Updated from reports to the Division of STD Prevention, NCHSTP.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending February 12, 2000, and February 13, 1999 (6th Week)

Reporting Area UNITED STATES NEW ENGLAND Maine N.H.	Cum. 2000 ^t	Cum.	Chiam				Escherichia coli O157:H7* NETSS PHLIS					
UNITED STATES NEW ENGLAND Maine	2000 ^s	2000 ^s			Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum
NEW ENGLAND	2.750	3,075	2000 39,345	1989 74,212	2000 93	1999	135	1999	2000	1999		
Maine	289	158	2,316	2,282	2	5	14	22	12	25		
	3	3	135	57	1	1	1	1				
18.	3	3	88	120	1	1	3	1	3	1		
Mass. 3.1.	234	122	1,161	989 258		2	3	13	2	13		
Conn.	42	19	868	816		1	6	7	6	11		
MID. ATLANTIC	795	486	438	8,524	9	21	20	6		2		
Upstate N.Y. N.Y. City	21 495	18 236	N	4,352	4	7	20	3		1		
V.J.	194	158	64	1,389			-	2		1		
Pa.	85	74	374	2,783	1	2	N	N				
E.N. CENTRAL Dhio	143	177	8,574 1,824	12,383	10	27	14	26 16	4	16		
nd. II.	26 64	25 77	1,324	1,194	3	2	1	4	1	3		
Mich.	19	22	2,130 2,329	3,145 2,146	2	3 2	6 3	2 4	i	2 2		
Nis.	9	16	967	1,473	-	16	N	N	1	3		
W.N. CENTRAL Minn.	49	114	1,862 506	4,398 947	2	7	30	23	20	14		
owa	7	22	101	167		1	7	6	9	9 2		
Mo. N. Dak.	15	73	686	1,800	2	4	18	2	7	1		
S. Dak.	1		112	265		*		2		1		
Nebr. Kana.	11	10	263 194	473 647		1	2	2 6	2	1		
S. ATLANTIC	588	845	8,229	16,929	11	7	16	12	9	7		
Del. Md.	15 92	13 81	338 639	316 1,695	i	2	6	î	1			
D.C.	22	8	302	N		2	-		Ü	Ú		
Va. W. Va.	41	54	867	1,786 274		*	3	4	2	2		
N.C.	27	68	2,111	2,504	2	1	4	2	1	2		
S.C. Ga.	36 97	56 110	669	3,409 3,364	3	i	1	1	3	1		
Fla.	256	445	2,662	3,582	5	1	2	3	2	1		
E.S. CENTRAL	140	156	3,951	3,990	3	2	5	12	.1	4		
Ky. Tenn.	20 36	15 62	950 1,168	782 1,607	-	1	2 2	3 6	1	U		
Ala. Miss.	50 35	30 46	1,102	1,363	3		1	2	:	2		
W.S. CENTRAL	276	530	731 3,236	238 9,450	4	4	4	2	ā	1 6		
Ark. La.	8	19 26	375	567 784	1		2	*		2		
Okla.	10	6	908	1,097	-				3	1		
Tex.	213	479	1,952	7,002	3	4	2	1	1	3		
MOUNTAIN Mont.	102	45	2,234	3,963	6	14	15 5	6	3	4		
ldaho	3	4	84	186	1	2	1					
Wyo. Colo.	34	26	432	81 781			4	1 2	î	1		
N. Mox.	8	4	94	614	1	8		-				
Ariz. Utah	22 12	4	916 319	1,591 231	2 2	A N	1	1	2	2		
Nev.	21	3	327	355		-	i					
PACIFIC Wash.	360	567 28	8,506 1,516	12,323 1,436	46 N	26 N	17	14	2	17		
Oreg.	11	15	374	615	1	3	3	8	1	6		
Calif. Alaska	299	509 5	6,373	9,696	45	23	11	6		6		
Hawaii	10	10	2-0	366			2	-				
Guam	-	1		50		-	N	N	U	U		
P.R. V.I.	77	92	113	U	2	Ü	:	ů.	U	Ü		
Amer. Samoa C.N.M.I.				Ŭ		ŭ		Ü	Ü	Ü		

N: Not notifiable

U: Unavailable

Cn.M.L.: Commonwealth of Northern Mariana Islands

Individual cases may be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health, Laboratory information System (PHLIS).

Updated monthly from reports to the Division of HiV/AIDS Prevention-Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention, last update January 30, 2000.

Chlamydia refers to genital infections caused by C. trachomatis. Totals reported to the Division of STD Prevention, NCHSTP.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending February 12, 2000, and February 13, 1999 (6th Week)

	Gone	orrhea	Heps C/N/	etitis A,NB	Legion	nellosis		TTIE BBSe
Reporting Area	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
INITED STATES	21,363	1999 41,501	2000 176	1999 367	2000 55	1999	2000	1999 459
EW ENGLAND	794	804		2	3	5	32	53
aine	8	7		-	2			
Н.	9	9 5	*	1	*	1	11	*
t. lass.	344	317	-	1	1	2	21	52
J.		77			-	1		-
onn.	432	389				-		1
ID. ATLANTIC pstate N.Y.	589 275	4,628 332	•	8	3 2	17	160 56	286 38
Y. City	2/5	2,048			-	4	1	12
J.	52	922	-	5	i	3		79
b.	262	1,326	_			8	103	157
N. CENTRAL	4,892 999	7,349 1,963	32	234	15 11	36 12	1	18 7
id.	596	743		-	2	1	*	*
lich.	1,077	2,362 1,578	30	75	2	6		1
lis.	592	703	30	156		7	Û	9
.N. CENTRAL	747	2,368	24	26	4	3	2	6
linn.	205	351	-	-	1		1	
lo.	31 324	1,413	24	24	1 2	2	1	1 2
. Dak.	-	7	-		-			1
, Dak. lebr.	8 91	24 221		i	5		*	
ens.	88	264		1		-	-	2
ATLANTIC	6,963	13,526	6	25	18	8	30	62
el.	184	187	:		1	1		3
ld. .C.	318 312	2,161 975	1	16	6	-	24	50
n.	971	1,504	*	2	2	2	*	-
/. Va. I.C.	1,963	2,337	3	1 5	N 1	N 2	1 3	
.C.	574	1,744		1	2	1		
a.	556 2.085	1,903 2,625	2	-	6	2	2	
S. CENTRAL	3,040	3,454		20	1	4	2	8
V.	426	460	32	2			:	
enn.	1,001	1.344	8	14	5	2 2		2
da. Aiss.	935 678	1,342 308	3 18	1 3	1	2	-	3
S. CENTRAL	1,786	5,429	35	5				
rk.	242	294	30		-	-	-	
a. Ikia.	458	859 597		2	*		-	
836.	1,088	3,679	36	2				
OUNTAIN	899	1,180	23	30	4	4	1	1
lont.		1	-			-		
laho /yo.	4 5	10	13	3 15	1	-		
olo.	410	213	4	3	2	1		
I. Mex. triz.	18 285	134 634	3	6 2	2	1	i	1
Itah	48	23	3	1	1	2		
lev.	129	162	-	-	-	-		
ACIFIC	1,863	2,763	24	17	7	4	12	25
/ash. reg.	289	247	2 5	2	1 N	Ň	1	
alif.	1,288	2,303	17	14	6	4	11	25
laska lawaii	29	70		-		-	Ň	Ñ
uam		12						14
R.	28	36	:				N	N
.1.		U		U	-	U		U
mer. Samoa		U		Ü		U		U

N: Not notifiable

U: Unavailable -: no reported cases

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending February 12, 2000, and February 13, 1999 (6th Week)

					Salmonellosis®						
	Mal			Animal		TSS		LIS			
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1990			
UNITED STATES	1,170	1,486	265	842	464	787	530	1,191			
NEW ENGLAND	31 2	32	12	36	6	9	11	23			
V.H.	1	2		4		-	-				
/t. Mass.	21	24	11	23	5	5	9	7			
R.I. Conn.	5	3 2	1	3 4	1	3	2	9 7			
MID. ATLANTIC	28	107	29	600	9	30	66	145			
Jpstate N.Y. I.Y. City	13	22 34	3 28	16	6	13	37	6 58			
V.J.	5	33	•	23	3	9	22 8	46 36			
E.N. CENTRAL	200	329	42	121	107	107	33	115			
Ohio nd.	14 18	127	1	9	9 50	10	11	37			
II.	68	111	5	96	14	28 57	17	9 51			
Mich. Nis.	96	36 44	34 2	10	23 11	7 6	3	16			
W.N. CENTRAL	57	76	31	66	6	30	23	27			
Minn. owa	12	10	12 7	14	2	1	13	16			
Mo. N. Dak.	25	54	8	46	4	27	8	9			
S. Dak. Nebr.	i	6	2	3		i	2	1			
Kans.		6	2	3		i	-	1			
S. ATLANTIC	97	147	15	38	176	322	83	118			
Md. D.C.	10	11 6	2	2	23 10	57		19			
√a.	9	5		3	17	32 21		4 9			
W. Va. N.C.	8	3	4	9	60	72	5 9	5 29			
S.C. Ga.	3 6	15	1 3	5 8	11	33	18 24	33 16			
Fla.	62	57	6	10	42	42	27	1			
E.S. CENTRAL Ky.	44 9	205	19 U	122 U	77	144	31	71			
Tenn.	19	149	17	114	52	63	4	16			
Ala. Miss.	11	22 14	2		14 8	44 20	27	7			
W.S. CENTRAL	84	211	63	299	42	94	11	234			
Ark. La.	18	15 11	10	11 18	3	10	8	8			
Okla. Tex.	66	122	52	9 261	12	24 56	3	220			
MOUNTAIN	142	101	33	57	18	16	17	30			
Mont. Idaho	15	2		i	-	-					
Wyo. Colo.	16	19	7	17	2	-	i	Ü			
N. Mex. Ariz.	17	10	6 17	6 25	14	16	3	4			
Utah Nev.	5	5	4	6		-	8 4	12			
PACIFIC	27 487	3 258	21	2 36	2 23	36	1 256	428			
Wash. Oreg.	57 65	4 7	2	18	4	1	21	12			
Calif.	359	239	10	9	19	32	226	10 383			
Alaska Hawaii	2 4	i		i		i	1 8	6 17			
Guam		2	U	U							
P.R. V.J.	:	8	Ü	Ü	18	32 U		Û			
Amer. Samoa C.N.M.I.		Ü	Ü	Ü		Ü		Ü			

N: Not notifiable U: Unavailable :- no reported cases
*Individual cases may be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public
Health Laboratory information System (PHLIS).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States,

-	Na	Shigel		LIŚ	Syr (Primary &	ohilis Secondary)	Tuber	culosis
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area JNITED STATES	1,170	1,466	265	1999 842	2000 464	787	530	1999 ¹
NEW ENGLAND	31	32	12	35	6	9	11	23
Maine	2			i		-		-
V.H. /t.		2		1		i	-	
Vlass. R.I.	21	24	11	23	5	5	9 2	7 9
Conn.	5	2	1	4	1	3		7
MID. ATLANTIC Upstate N.Y.	28 13	107	29	69 16	9	30	65	145
N.Y. City	10	34	26	30	6	13	37	58
N.J. Pa.	6	33 18		23	3	9	6	46 36
E.N. CENTRAL	200	329	42	121	107	107	33	115
Ohio ind.	14 18	127	1 5	9	9	10 28	11 2	37
II.	68	111	*	98	14	57	17	51
Mich. Wis.	96 4	36 44	34	10	23	7 5	3	15
W.N. CENTRAL	57	76	31	66	6	30	23	27
Minn. Iowa	12 12	10	12	14	2	1	13	16
Mo. N. Dak.	25	54	8	46	4	27	8	9
S. Dak.				-				î
Nebr. Kans.	8	6	2 2	3		1	2	1
S. ATLANTIC	97	147	15	38	176	322	83	118
Del. Md.	10	11	2	1 2 U	23	57		2 19
D.C. Va.	9	6 5	Ü	3	10 17	32 21		4 9
W. Vn.		3				1	5	5
N.C. S.C.	8	38 15	4	9 5	80 11	72 33	9	29 33
Ga. Fla.	5 62	8 57	3 5	10	12	63	24 27	16
E.S. CENTRAL	44	206	19	122	77	144	31	71
Ky. Tenn.	9	149	17	114	3 52	63	4	6
Ala.	5	22	-	8	14	44	27	42
Miss. W.S. CENTRAL	11	14 211	63	299	8	20 94	11	234
Ark.	18	15		11	3	10	8	8
La. Okla.	-	63	10	18	27	24	3	Ü
Tex.	66	122	52	261	12	56		220
MOUNTAIN Mont.	142	101	33	57	18	16	17	30
ldaho Wyo.	15	2		1				
Colo.	16	19	7	17	2		1	Ü
N. Mex. Ariz.	17 62	10	5 17	6 25	14	16	3 8	12
Utah Nev.	5 27	5 3	4	6 2	2		4	8
PACIFIC	487	258	21	36	23	35	256	428
Wash.	57	4 7	2	18 9	4	1	21	12
Oreg. Calif.	65 359	239			19	32	226	383
Alaska Hawaii	2 4	8	-	8	-	i	1 8	17
Guam		2	U	U				
P.R. V.I.		6	U	U	16	32 U	:	ú
Amer. Samoa		Ü	U	Ü		U		U
C.N.M.I.	Utille	U	U	U	•	U		U

N: Not notifiable

I: Unavailable

In or reported cases

Individual cases may be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

I'cumulative reports of provisional tuberculosis cases for 1999 are unavailable ("U") for some areas using the Tuberculosis Information System (TIMS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending February 12, 2000, and February 13, 1999 (6th Week)

	H. influ		Н	epatitis (Vi	ral), by typ					ies (Rubec		
1	inva		A		В		Indige		Impo		Tota	
Reporting Area	Cum. 2000 ¹	Cum. 1909	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	2000	Cum. 2000	2000	Cum. 2000	Cum. 2000	Cum. 1999
NITED STATES	97	123	1,097	1,724	403	541	-	1			1	13
EW ENGLAND	6	9	18	27	6	16			*			
taine I.H.	1	i	1	2 2	1 3	2				-		
۴.	1	2	1		2		-		-			-
fass.	4	6	3	10	-	6 2	1		-			-
onn.	-	-	9	13		6				*	-	
AID. ATLANTIC	12	19	38	115	23	82			-		-	
Ipstate N.Y. I.Y. City	10	9	26 12	12 47	17	12 23	-	-	*	-		-
I.J.	1	4		24		14	U		U			
а.	1	*	*	32	*	33				*	*	*
N. CENTRAL	13	24	126	465 77	58 13	14		1	-	*	1	
Ohio nd.	8 2	1	52	9	1	4	-		-	-		
II.	2	12	10	93	44	41		-			i	
Aich. Vis.	1	-	1	278	-	4	-	1				
W.N. CENTRAL	2	5	109	96	17	28						
Minn.		-	12		*							
owa Mo.	î	1 2	11 80	7	14	18	:					-
N. Dak. S. Dak.		*			-		U		U			-
S. Dak. Nebr.	1	1	6	9	î	6	U		Ü	*	*	*
Cans.	-	1		6		2	U		U			~
S. ATLANTIC	35	23	98	131	65	75	~	*	-	*		-
Del. Md.	17	17	18	49	17	32						-
D.C.				7		*						
∕a. W. Va.	8	1	16 7	9	16	6				-		*
N.C.	3	2	20	19	11	26						
S.C. Ga.	1 4	2	2 4	46	1	8						-
Fla.	1		31		21	-		-				-
E.S. CENTRAL	3	10	51	54	31	46						
Ky.	2	2 4	2 15	9	23	2 23						-
Tenn. Ala.	1	3	8	16	2	11		-			-	-
Miss.		1	26	9	5	8		-				-
W.S. CENTRAL	-	6	133	172	6	44		-				2
Ark. La.			11	3	6	7	Ü		Ü			
Okla.		5		61	*	10					-	
Tex.	*	1	122	107		26		*		-	-	2
MOUNTAIN Mont.	16	16	85	181	42	57						-
Idaho	1	1	3	4	3	4						
Wyo. Colo.	5	1	26	40	11	13		-	*	*	*	
N. Mex.	5	3	9	5	12	20						-
Ariz.	6	6	31	99 12	14	9 5		*	*			-
Utah Nev.		3	8 7	19	1	6				-		
PACIFIC	8	11	430	484	155	132			-		_	11
Wash.	2 2		3 27	8	1	1						2
Oreg. Calif.	2	3 7	406	22 451	13 138	121	-			-		8
Alaska	1	1	3	2	2	2						
Hawaii	3			1	1	1						
Guam P.R.	:	:		2 7	-	14	U	-	U	-		
V.I.		U		Ú		U	U	-	Ü	-	,	U
Amer. Samoa C.N.M.I.	:	U		Ü		U	Ü		U	*	-	Ü

N: Not notifiable U: Unavailable -: no reported cases
*For imported measles, cases include only those resulting from importation from other countries.
**Of 26 cases among children aged c5 years, serotype was reported for 10 and of those, 2 were type b.

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending February 12, 2000, and February 13, 1999 (6th Week)

		ococcal ease		Mumps			Pertussis		Rubella			
Reporting Area	Cum. 2000	Cum. 1909	2000	Cum. 2000	Cum. 1999	2000	Cum.	Cum.		Cum.	Cum	
NITED STATES	265	238	11	35	43	33	337	1999 425	2000	2000	1999	
EW ENGLAND	14	18			3	5	72	62	1	1	1	
aine	1	2	-			2	4					
.Н.	1	2	-		1	2	20 24	3 7	1	1		
288.	7	13	-		2		23	52		*	1	
l. onn.	1 4				-	1	i		-	*		
ID. ATLANTIC	19	31		2	5	7	24	21				
ostate N.Y.	6	4		î		7	22	11				
Y. City J.	3	13	ú	*	2	ū	*	7	11	*		
1.	6	6	-	1	3		2	2	u	:		
N. CENTRAL	32	35	-	1	2	8	96	61				
hio	9 7	15		-	1	6	89	41				
d.	4	13	-	:	1	2	3	6				
ich.	11	2 2	*	1			3	5	*			
is.	1						*	8			-	
.N. CENTRAL	30	24	3	6	1	*	7	11	*			
wa	3	4		1	1		3	4				
o. Dak.	26	12	Ü	1	-	Ü	1	1	ű			
Dak.		3	U	-	-	ŭ		1	Ü			
ebr. ans.		1 4	2	4		ű	*	6	Ü		-	
ATLANTIC	52	26	1	4	5		-		U			
al.	502	1		4	5	2	23	42	-			
d.	4	6		1	1	1	9	18	-			
.C.	9	2			-	-	1	6				
.Va.	.1	1		*	2	*		-				
.C. C.	11	5	1	3	1 2	1	9	16 2	*			
a. a.	7	4				-						
					1	*	~					
S. CENTRAL	10	23		1		-	7 3	12	-	-		
inn.	3	8			*		1	4				
la. liss.	5	9		1			3	6	-			
S. CENTRAL	1	17			9		1	16				
rk.	1	3					1	2				
i. kia.		5	U		i	U			U			
IX.		3			8	*		12				
OUNTAIN	14	28		2	3	9	95	91		1		
ont.	2	ã			-					-		
aho yo.	2	1				2	15	44				
olo.	1	8			1	5	52	14				
. Mex.	6	7	N	N	N	1	16	7 9				
tah	3	3		-	1	-	3	15		1		
ev.		1	-	2	1	1	1	1				
CIFIC ash.	93	37	7	19	15	2	12 2	109	*			
reg. alif.	13	8	N	N	N	2	8	3	-			
slif. Iaska	75	18	7	19	11	-	2	100	*	-		
swaii	1	4			3		-	4	-			
uam	-		U			U			U			
R.	*	ú	U		Ü	U		12	U	-		
I. mer. Samoa	:	U	Ü	2	Ü	Ü		U	Ü	-	U	
N.M.I.		Ü	Ü		Ü	Ŭ	-	ŭ	ŭ		ŭ	

TABLE IV. Deaths in 122 U.S. cities,* week ending February 12, 2000 (6th Week)

		All Car	uses, By	Age (Y	ears)		PMI			All Cau	ses, By	Age (Y	(ears)		PBJ
Reporting Area	All Ages	≥66	45-64	25-44	1-24	<1	Total	Reporting Area	Ali Ages	≥65	45-64	25-44	1-24	<1	Tota
NEW ENGLAND Boston, Mass. Sridgeport, Conn. Lambridge, Mass. all River, Mass. All River, Mass. Jynn, Mass. New Bedford, Ma New Haven, Conn Providence, R.I. Somerville, Mass. Springfiel Mass.	. 25 33 U 31 17 88. 30 . 43 74	473 95 33 21 32 U 22 14 26 59 6	105 33 9 4 1 U 7 2 4 10 9 2 12	29 2 6	4 2 U	11 2 1 · · · · · · · · · · · · · · · · ·	77 15 8 4 4 U 3 2 2 2 6 4	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ge. St. Petersburg, I Tampa, Fla. Washington, D.	103 67 82 68 Fla. 50 192 C. 190	813 U 124 76 91 66 50 57 57 144 116 3	246 U 42 22 37 25 10 13 10 6 37 39 5	106 U 27 2 18 7 1 7 3 3 6 16 15	45 U 11 3 5 5 1 2 2 3 13	29 U 1 2 2 1 6 4 3 2 2 6	114
Vaterbury, Conn. Vorcester, Mass. AID. ATLANTIC Ilbany, N.Y. Illentown, Ps. Juffalo, N.Y. Juffalo, N.J. Jilizabeth, N.J. Irle, Ps. §	36 74 2,779 49 U 190 28 23 67	30 58 1,970 33 U 148 17 14 56	523 12 U 28 7 6	176 176 1 U 9 2	54 1 U	1 3 56 2 U 5 1	6 16 186 3 U 21 1	E.S. CENTRAL Birmingham, Al Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphia, Tenn Mobile, Ala. Montgomery, A Nashville, Tenn.	1,106 a. 214 ann. 108 96 98 262 141	772 162 77 66 37 165 99 59	235 39 23 19 18 64 31 8 33	53 6 5 6 1 17 6 2	23 5 1 1 7 5	23 2 2 4 1 9	131 2 3 1 1 1 1 1
lersay City, N.J. New York City, N.J. Paterson, N.J. Philadelphia, Pa. Pittsburgh, Pa. Seading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa. S. Fyracuse, N.Y. Jitica, N.Y. Yonkars, N.Y.	36 23 488 60 36 131	925 18 12 365 44 27 101 23 28 65 30	10 265 10 5 88 11 6 20 4 9 12 13	3 101 7 5 19 2 4 7 3 1 5 5	1 29 1 16 2 2	2 24 2 1 10 1 1 3 	42 3 3 37 10 2 12 3 19 11 8 0	W.S. CENTRAL Austin, Tex. Baton Rouge, Le Corpus Christi, Dallas, Tex. Fl. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, Le San Antonio, Te Shreveport, Le Tulisa, Okla.	Tex. 54 U 90 157 389 86	993 62 87 40 U 36 119 245 51 48 166 35	318 16 29 10 U 11 26 92 22 18 44 18 32	113 6 12 3 U 1 9 39 8 7 16 1	26 U 1 1 10 4 8 1	25 1 3 1 U 1 2 3 5 3 5 1	14
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	2,329 98 52 438 122 123 222 161 216 63 94	1,645 43 40 267 88 81 167 127 119 50 69	418 11 9 93 20 31 36 25 51	163 2 2 46 7 7 15 6 31	41 9 5 1 1 3 5	22 1 24 2 3 1 1 10 1 2	6	MOUNTAIN Albuquerque, N Boise, Idaho Colo. Springs, (Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, L Tucson, Ariz.	39 Colo. 51 124 214 28 174 25	743 86 31 40 82 143 24 119 19 70 129	206 25 7 8 23 54 4 34 5 14 32	72 7 1 2 13 14 13 1 10 11	19	14 2 4 1	•
Gary, Ind. Grand Rapids, Mi Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngatown, Oh	159 56 118 53 64 46 129	18 42 115 42 88 44 46 39 96	8 8 18 7 10 4 20 8	1 5 7 3 9 1 6 2 6	1 3 5 1 2	3 6 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 12 25 2 14 2 10 6	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Haw Long Beach, Ca Los Angeles, Ca Pasadena, Calif. Portland, Oreg. Sacramento, Ca	lif. 90 lif. 289 18 115	1,063 6 93 12 61 70 219 13 83	247 2 33 4 10 12 46 5 20 U	72 1 7 2 1 5 14	25 2 2 1 5	21 1 2 2 2 6	16
W.N. CENTRAL Des Moines, low Duluth, Minn Kansas City, Kans Kansas City, Mo. Lincoln, Nebr. Minneapolis, Mir Omaha, Nebr. St. Louis, Mo. St. Paul, Minn.	118 42	74 32 198 76 85	196 23 U 7 30 7 35 16 34	6 6 6 2 13 7	32 3 U 2 3 1 4 1 4 4	22 4 U 1 5 - 2 - 5 2 3	29 U 4 14 7 34 18 2	San Diego, Cali San Francisco, C San Jose, Calif Santa Cruz, Cal Seattle, Wash. Spokane, Wash Tacoma, Wash. TOTAL	f. 177 alif. U . 190 f. 23	124 U 144 20 107 43 66	30 U 32 3 26 8 16	14 U 7 8 2 4	5 U 4 - 2 1 269	263	1,3

U: Unavailable :no reported cases

*Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of 100,000 or more.

A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.

Pneumonia and influenza.

*Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.

*Total includes unknown ages.

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